



Antibacterial Activity of Ag-Cu-O Nanocomposite - Thermal Reduction Method

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ABSTRACT

The development of nanocomposite with antimicrobial property and stability is of great interest nowadays. The incorporation of silver, whose microbial activity is well known in the copper oxide matrix, will result in an amazing property. This new nanocomposite a potential material in the medical, food packaging, water treatment and pharmaceutical fields. Copper and silver are abundant in nature and cost-effective too. To synthesis high purity Ag-Cu-O nanocomposite novel thermal degradation method is used in the present paper. Structure, surface and optical characteristics are studied in detail. The microbial activity of the synthesized nanocomposite in Gram-positive and negative bacteria are studied by disc method and compared with the existing reports.

Keywords: Antimicrobial activity; Nanocomposites; Silver copper oxide; Structural analysis; Surface morphology.

1. INTRODUCTION

Nanotechnology is playing a very crucial role in many fields, including agriculture, medicine, environmental science and textile and food industries. Particularly in the field of medicine, metal nanoparticles like Ag, Au, Cu, Ti and Pt nanoparticles are showing good antibacterial and antifungal properties (Jose *et al.* 2005; Pinto *et al.* 2012). In recent times the interest is increasing in the field of nanocomposites where two or more nanoparticles will combine to result in superior properties. The physical and chemical properties of nanocomposites are different from their parent nanoparticles. Improved stability and hardness are the main characteristics of metal nanocomposites. In this context, the development of metal nanocomposite with antimicrobial activity is interesting because of its size and shape-dependent properties. Among all the metal nanoparticles, Ag and Cu have shown appreciable antibacterial activity in many bacterial strains. These nanoparticles are found to be used in various fields such as biomedical equipment, water treatment and food processing (Ruparelia *et al.* 2008; Llorens *et al.* 2012; Mohammad J. Hajipour, 2012). The cationic release is the main advantage of silver nanoparticle's strong antimicrobial and fungal activity (Asha rani, 2009). It is always attentive towards the search for alternatives nanoparticle, which gives comparable results. Many researchers have reported that the copper nanoparticles are resulting analogous effects in many pathogen microorganisms (Cady *et al.* 2011; Turnlund, 1998;

Yoon *et al.* 2007). In several instances, the metal oxide nanocomposites demonstrated better performance over the mono metal oxides (Gitashree Darabdhara *et al.* 2017; Tan *et al.* 2014; Jiang *et al.* 2012; Wu *et al.* 2015).

Based on this, the present paper discusses the silver-copper oxide nanocomposite synthesis, and its biological activity is studied by *in situ* method using Gram-positive and Gram-negative microorganisms. The purity of the material plays a vital role in biological properties. In the present work, the thermal degradation method was used to synthesis silver-copper oxide (Ag-Cu-O) nanocomposite.

2. EXPERIMENTAL TECHNIQUES

The thermal degradation method is known for its controlled synthesis of metal nanoparticles. This is a unique method where the purity of the synthesized nanoparticles is more than 99.9 % as this method won't require any organic solvents to clean the resultant product. Also, the morphology and size of the samples can be easily controlled by varying the temperature without any capping agent. In the present work Cu:Ag ratio is to 8:2. All substances used were of analytical grade and 99.99 % purity. The solvents used were distilled before use. 3-hydroxy 2- naphthoic acid and aminoguanidine were purchased from Aldrich, and all other chemicals used were commercially available grades. The synthesis of Ag-Cu-O nanocomposite is as follows.

2.1 Preparation of Copper-Silver Oxide Complex

A known weight of 3-Hydroxy 2-Napthoic acid (0.1881grams, 0.001 mol) was dissolved in 20 ml of double-distilled water. Aminoguanidine (0.544grams, 0.004 mol) was added to the above solution while stirring. The solution became clear, and the pH (7) was measured. Copper nitrate solution (0.1876 g) and silver nitrate solution (0.1688 g) was dissolved in 20 ml of water. The solution was poured into the metal solution, and the resulting solution was concentrated for five hours on a water bath. A polycrystalline substance obtained after two days was washed with distilled alcohol and dried in desiccators.

2.2 Synthesis of Metal Nano Composite and Nano Metal Oxides

Ag-Cu-O nanocomposite was synthesized by the thermal decomposition method. The complexes were weighed (12 g) and placed in a muffle furnace at a temperature of 38 °C and raised steadily up to 800 °C at a heating rate of 10 °C per minute. When the temperature has reached 800 °C, it was maintained for 6 hours. The metal oxides and composites were prepared in a similar manner using the same procedure.

2.3 Physico-Chemical Experiment Techniques

The synthesized metal nanocomposites were inferred through physicochemical techniques like UV-DRS, FTIR, XRD and SEM with EDAX studies. The infrared spectra of the pure nano metal oxides and their composite were recorded on the IR Affinity – 1CE Schimadzu model spectrometer in the range 4000 – 400 cm. UV visible spectra of pure metal oxides and their composite were recorded on Lab India UV 3000+ UV-Vis spectrometer. XRD of the pure nano metal oxides and their composite were recorded on Bruker Advanced D8 X-ray diffractometer. SEM with EDAX was observed using Joel JSM-6390 LV SEM at a 3-12 KV at different magnifications. Antibacterial studies were carried out in *Staphylococcus aureus*, *Escherichia coli*, *Bacillus subtilis* bacterial stains by in situ method.

3. RESULTS & DISCUSSION

3.1 Powder X-ray diffraction

The diffractogram was recorded using the synthesized recorded at room temperature. The diffraction pattern was recorded from 10°- 90° at the scan rate of 0.2 °/min. Figure 1. shows the XRD pattern Ag-Cu-O nanocomposite. The sharp peaks in the XRD pattern revealed that the formed materials are crystalline in nature. The synthesised metal oxide XRD peaks are in accordance with JCPDS card number 89-3722 (Ag₂O) and 65-2309 (CuO) (Gitashree *et al.* 2017). The

nanocomposite peaks are well-matched with the above results and indexed according to that. This confirms the formation of silver-copper oxide composite. The absence of impurity peaks in XRD pattern revealed the high purity of the synthesised nanoparticles. The structure of the nanocomposite is faced centred cubic. Using the Scherrer formula, the average crystallite size was calculated, and it was about 32 nm.

3.2 Fourier Transform Infrared spectroscopy (FTIR)

Fig. 2. shows the FT-IR spectrum of the nanocomposite. ATR mode was used to record the spectrum at room temperature. The spectrum displays two characteristics broad band centred at 3517 and 1521 cm⁻¹ which are referring to the stretching and bending mode vibrations. The band at 739 cm⁻¹ corresponds to the vibration of Metal-oxide stretching mode (Devi *et al.* 2010; Ahmed, 2012). The strong band at 425 cm⁻¹ corresponds to the vibration of Ag-Cu bond. It could be seen from figure 3.1. that the broad absorption band centred at 3450 cm⁻¹ is attributed to the band OH stretching vibrations due to the fact that the calcinated powders tend to physically absorb water. The sample contains different proportion of copper oxide exhibit some small bands at 1219 cm⁻¹ which is attributed to copper oxide.

3.2 Scanning Electron Microscopy (SEM)

SEM is a very effective tool to study the surface morphology of nanomaterials. A micrograph of the Ag-Cu-O nanocomposite is shown in Figure. Well, disperse the plate-like structure, and the tubes are visible on the surface of the synthesized material. Copper forms a plate-like structure, whereas silver prefers to have a tube-like structure. This was also confirmed with the EDX analysis, which is shown in Fig. 3 and the inset. Elemental mapping was carried out individually at the plate and the tube and found that the wt % of Cu and Ag is high, respectively, in that structures (Fig. 4).

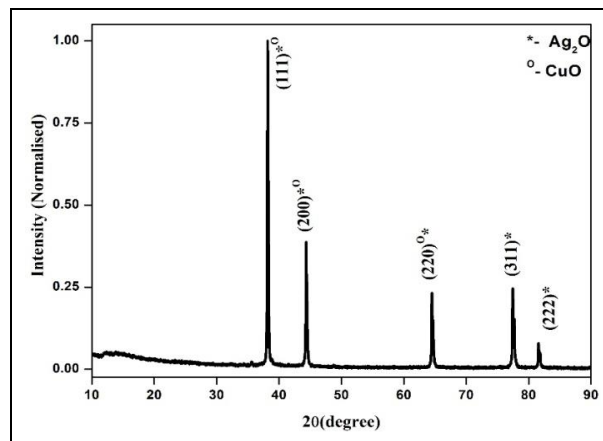


Fig. 1: XRD Spectrum of Ag-Cu-O

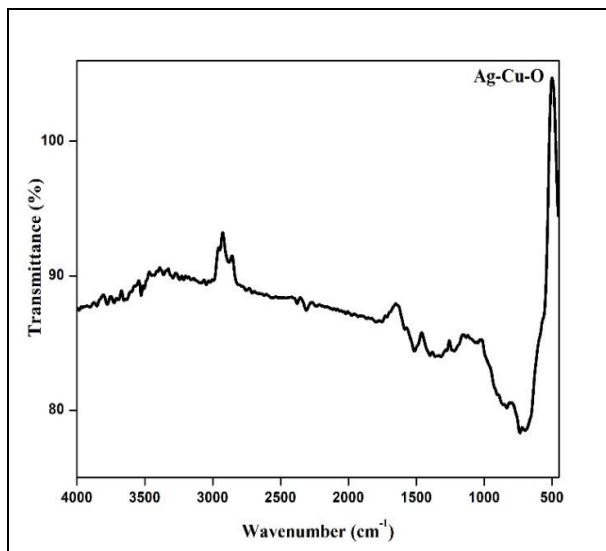


Fig. 2: FTIR Spectrum of Ag-Cu-O

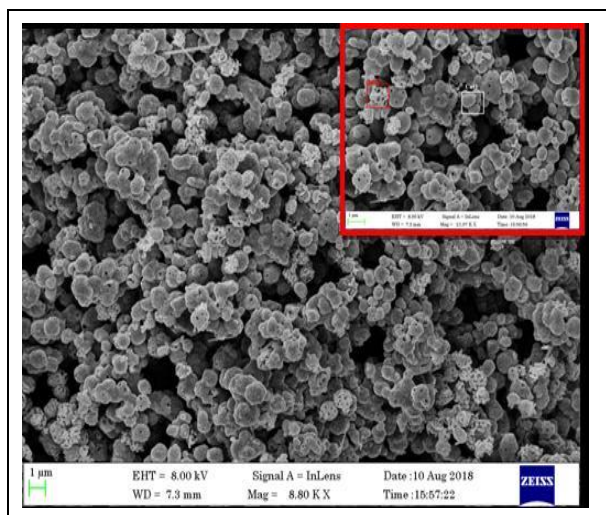


Fig. 3: SEM of Ag-Cu-O (Inset: Zoomed View)

3.4 UV - Visible Diffuse Reflectance Spectroscopy

UV-Vis DRS is a unique tool to measure the reflectance of nanomaterials. Ag-Cu-O absorption spectrum is shown in Figure 5. The broad absorption at 320 nm is a significant plasmon band of Ag and Cu metals. $\pi \rightarrow \pi^*$ energy transfer mechanism is attested from this absorption. Other than this absorption, the synthesized nanocomposite is transparent in the visible region.

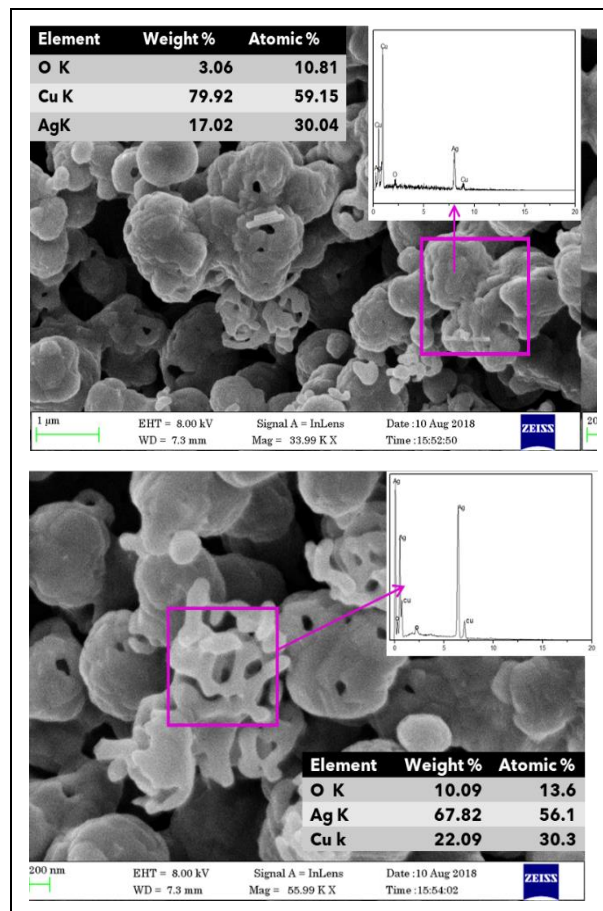


Fig. 4: Elemental Mapping of the Plate and the Tube-like Structures

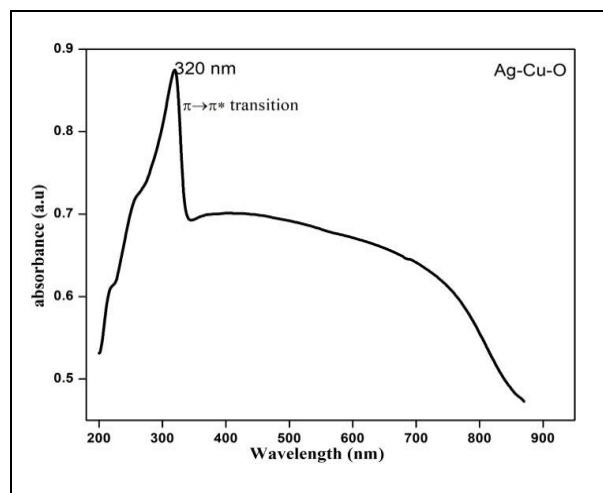


Fig. 5: UV-DRS Spectra of Ag-Cu-O

3.5 Antibacterial activity

Antibacterial activity of in situ synthesized Ag-Cu-O nanocomposite was tested by standard disc method using *Staphylococcus aureus*, *Escherichia coli*, and *Bacillus subtilis*.

3.5.1 Preparation of Inoculum

The inoculums for the experiment were prepared in fresh Nutrient broth from preserved slant culture. The inoculums were standardized by adjusting the turbidity of the culture to that of McFarland standards. The turbidity of the culture may be adjusted by the addition of sterile saline or broth or by further incubation to get the required turbidity. All the materials used, like cotton swabs, forceps and etc., are sterilized using alcohol.

3.5.2 Disc Method - measurement of the zone of inhibition

The standardized inoculums are inoculated in the plates prepared earlier (aseptically) by dipping a sterile in the inoculums removing the excess of inoculums by passing by pressing and rotating the swab firmly against the side of the culture tube above the level of the liquid, and finally streaking the swab all over the surface of the medium 3 times rotating the plate through an angle of 60 °C after each application. Finally, pass the swab around the edge of the agar surface. Leave the inoculums to dry at room temperature with the lid closed. Each Petri dish is divided into 2 parts, in one part prepared disc Ag-Cu-O (100 µg) disc (discs are soaked overnight in sample solution) and other part standards Ciprofloxacin 10 µg, are placed in the plate with the help of sterile forceps. Then Petri dishes are placed in the

refrigerator at 4 °C or at room temperature for 1 hour for diffusion. Incubate at 37 °C for 24 hours. Finally, observed the zone of inhibition produced by different samples and was measured using a scale and recorded the average of two diameters of each zone of inhibition. The results are incorporated in Figure 6 and the in table 1. From the results, it is obvious that the synthesised composite has antibacterial activity even without the presence of any antibodies. The better activity was observed in Gram-positive bacteria's.

Table 1. Antimicrobial Activity of Synthesized Ag-Cu-O Nanocomposites

S. No.	Organisms	Zone of Inhibition (mm)	
		std	Ag-Cu-O
1	<i>Staphylococcus aureus</i>	45	18
2	<i>Bacillus subtilis</i>	42	18
3	<i>Escherichia coli</i>	40	15

4. CONCLUSION

A unique thermal degradation method is employed to synthesis Ag-Cu-O nanocomposite. This method resulted in high purity FCC structures Ag-Cu-O, and this was confirmed from the XRD analysis. The presence of Ag and Cu was attested by the strong absorption band in FTIR spectrum. Plasmon transfer mechanism is evidenced by the deep absorbance in UV spectrum. The prepared nanoparticles exhibited good antibacterial property in gram-positive and negative bacteria which shows this the synthesized particles are suitable for wound dressing materials and for food packaging materials.

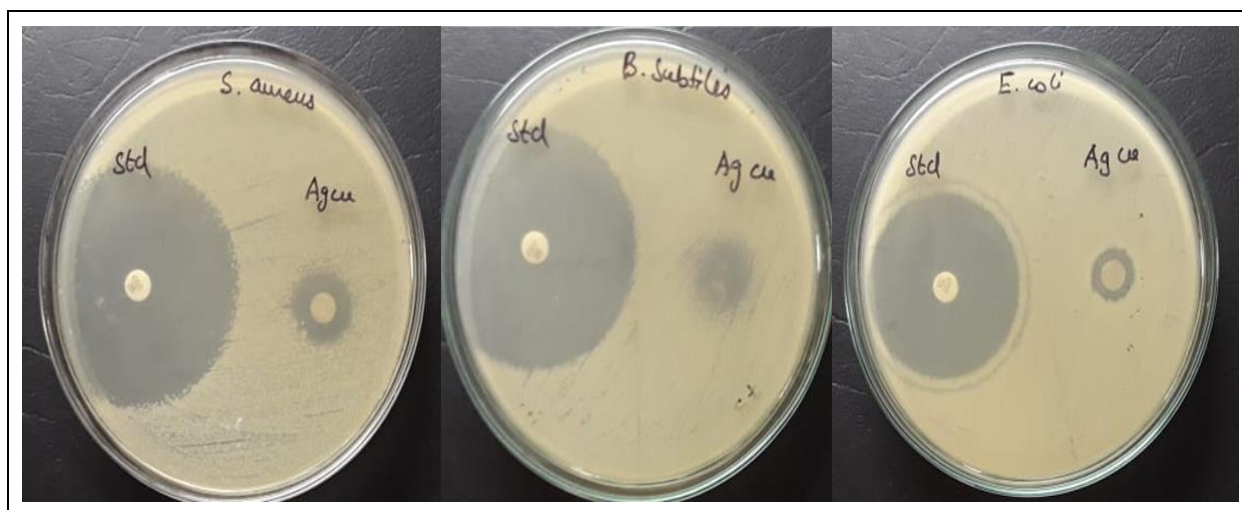


Fig. 6: Antimicrobial activity of Ag-Cu-O

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